

8EHQ-0601-13829



DuPont Haskell Laboratory

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for Toxicology and Industrial Medicine
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June 5, 2001



8EHQ-96-13829

Via Federal Express

Document Processing Center (7407)
Room G99 East Tower
Attention: 8(e) Coordinator
Office of Pollution, Prevention, and Toxics
U.S. Environmental Protection Agency
401 M Street SW
Washington, DC 20460-0001

Contain NO CBI



89010000210

Dear 8(e) Coordinator:

8EHQ-1296-13829

Mixture of 1, n-diiodoperfluoroalkanes (n = 4-10)

This letter is to inform you of the results of an acute inhalation toxicity study that was recently conducted with the above referenced test material.

Five groups of 6 male rats (10-12 weeks old) each were exposed (4-hour exposure) whole-body to aerosol/vapor concentrations of approximately 0.74, 3.7, 6.6, 12, or 19 mg/L of the test material. Two control groups of 6 male rats (10-12 weeks old) each were exposed to air only. One control group was fed *ad libitum* and the other was feed-restricted. The purpose for the two control groups in this study was to distinguish primary compound-related effects on target organs from those arising secondarily to anticipated body weight losses. The 6 rats exposed to 3.7 mg/L and the 12 control rats were evaluated for gross and histopathological changes after 3-day and 14-day recovery periods (3 rats per recovery period).

No deaths occurred at any of the concentrations tested. During and/or immediately following the 12 and 19 mg/L exposures, rats exhibited a diminished response to sound stimulus. Rats exposed to 19 mg/L also exhibited weakness immediately following the exposure. Following 14 days of recovery, rats exposed to 3.7, 6.6, 12, and 19 mg/L had overall mean body weight losses of 16%, 15%, 21%, and 33%, respectively.

Compound-related gross and histopathological effects were seen in the testes and liver of rats exposed to 3.7 mg/L following a 14-day recovery period. Gross observations included discolored/mottled livers and small seminal vesicles. Microscopically, rats had minimal to mild testicular degeneration/atrophy and fatty change in the liver. No gross or histopathological effects were observed in rats exposed to 3.7 mg/L following a 3-day recovery period. Rats exposed to 3.7 mg/L had increased liver weights following both a 3- and 14-day recovery period.

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Under these experimental conditions, the findings described above appear to be reportable, based upon guidance given in the EPA TSCA Section 8(e) Reporting Guide (June 1991).

Sincerely,

A handwritten signature in black ink that reads "A. Michael Kaplan". The signature is fluid and cursive, with a long horizontal stroke extending to the right.

A. Michael Kaplan, Ph.D.
Director - Regulatory Affairs

AMK/AJO:clp
(302) 366-5260